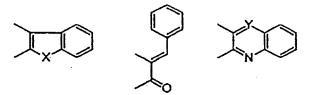
In the Claims

- 1.-10. (Cancelled)
- 11. (Currently Amended) A method of treating nausea and vomiting caused by a μ opioid receptor agonist compound comprising administering a therapeutically effective
 amount of an δ -opioid receptor antagonist agent comprising a morphinan derivative
 represented by general formula (I):

or a pharmacologically acceptable acid addition salt thereof as an active ingredient,

where R¹ represents a hydrogen atom, an alkyl group having 1 to 5 carbon atoms, a cycloalkylalkyl group having 4 to 7 carbon atoms, a cycloalkenylalkyl group having 5 to 7 carbon atoms, an aryl group having 6 to 12 carbon atoms, an aralkyl group having 7 to 13 carbon atoms, an alkenyl group having 3 to 7 carbon atoms, a furanylalkyl group (where the alkyl moiety has 1 to 5 carbon atoms), or a thiophenylalkyl group (where the alkyl moiety has 1 to 5 carbon atoms); R² and R³ are mutually independent and represent a hydrogen atom, a hydroxy group, an alkoxy group having 1 to 5 carbon atoms, an alkenyloxy group having 3 to 5 carbon atoms, an aralkyloxy group having 7 to 16 carbon atoms, an arylalkenyloxy group having 7 to 16 carbon atoms, an alkanoyloxy group having 2 to 6 carbon atoms, an alkenoyloxy group having 4 to 6 carbon atoms, an arylalkanoyloxy group having 7 to 16 carbon atoms, or an alkyloxyalkoxy group having 2 to 10 carbon atoms; R⁴ and R⁵ together form an -O-, -S-, or -CH₂- bond, or are mutually independent and R⁴ represents a hydrogen atom, a hydroxy group, an alkoxy group having 1 to 5 carbon atoms, or an alkanoyloxy group

having 2 to 6 carbon atoms and R⁵ represents a hydrogen atom; R⁶ represents a hydrogen atom, an alkyl group having 1 to 5 carbon atoms, an alkenyl group having 2 to 6 carbon atoms, an arylalkyl group having 7 to 16 carbon atoms, an arylalkenyl group having 7 to 16 carbon atoms, a hydroxyalkyl group having 1 to 5 carbon atoms, an alkoxyalkyl group having 2 to 12 carbon atoms, a COOH-group, or an alkoxycarbonyl group having 2 to 6 carbon atoms; and -Q-moiety represents a group as follows:



(where these structures may have one or more substituents selected from the group consisting of a fluorine atom, a chlorine atom, a bromine atom, an iodine atom, a nitro group, an alkyl group having 1 to 5 carbon atoms, a hydroxyl group, an oxo group, an alkoxy group having 1 to 5 carbon atoms, a trifluoromethyl group, a trifluoromethoxy group, a cyano group, a phenyl group, a hydroxyalkyl group having 1 to 5 carbon atoms, an isothiocyanato group, SR⁸, SOR⁸, SOR⁸, (CH₂), OR⁸, (CH₂), COOR⁸, SOONR⁹R¹⁰, CONR⁹R¹⁰, (CH₂), NR⁹R¹⁰, and (CH₂), N (R⁹) COR¹⁰ (where r is an integer from 0 to 5, R⁸ represents an alkyl group having 1 to 5 carbon atoms, R⁹ and R¹⁰ are mutually independent and represent a hydrogen atom, an alkyl group having 1 to 5 carbon atoms, or a cycloalkylalkyl group having 4 to 7 carbon atoms), and where X represents an oxygen atom, sulfur atom, a CH=CH, or NR⁷ group (where R⁷ represents a hydrogen atom, an alkyl group having 1 to 5 carbon atoms, an alkenyl group having 3 to 5 carbon atoms, an arylcarbonyl group having 7 to 13 carbon atoms, an alkylsulfonyl group having 1 to 5 carbon atoms, an arylsulfonyl group having 6 to 12 carbon atoms, an aralkylsulfonyl group having 7 to 16 carbon atoms, an arylalkenyl group having 7 to 16 carbon atoms, an arylalkenyl group having 2 to 6 carbon

atoms); and Y represents a nitrogen atom or a CH group to a mammal.

12. (Previously Presented) The method according to claim 11, wherein the -Q-moiety in general formula (I) represents a group:

(where X is as defined above and the group may have the substituents above).

- 13. (Cancelled)
- 14. (Previously Presented) The method according to claim 11, wherein R⁴ and R⁵ in general formula (I) together form an -O- bond.
 - 15. (Cancelled)
- 16. (Currently Amended) The method according to claim 11, wherein the μ-opioid receptor agonist compound is morphine.
 - 17.-22. (Cancelled)